

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

6900-255 M1S

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR

09/622781

INTERNATIONAL APPLICATION NO.
PCT/CA99/00155INTERNATIONAL FILING DATE
22 February 1999PRIORITY DATE CLAIMED
23 February 1998

TITLE OF INVENTION

LIGNIN BASED COLLOIDAL COMPOSITIONS

APPLICANT(S) FOR DO/EO/US

Zhi-Zhong LIANG

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☐ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c) (2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☐ A copy of the International Search Report (PCT/ISA/210).
8. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
9. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3))
10. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)). - unsigned copy
11. ☒ A copy of the International Preliminary Examination Report (PCT/IPEA/409).
12. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).

Items 13 to 20 below concern document(s) or information included:

13. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
14. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
15. ☒ A **FIRST** preliminary amendment.
16. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
17. ☐ A substitute specification.
18. ☐ A change of power of attorney and/or address letter.
19. ☐ Certificate of Mailing by Express Mail
20. ☐ Other items or information:

U.S. APPLICATION NO. (IF KNOWN) SEE 37 CFR

097622781

INTERNATIONAL APPLICATION NO

PCT/CA99/00155

ATTORNEY'S DOCKET NUMBER

6900-255 MIS

21. The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :

- ☐ Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO \$970.00
- ☒ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$840.00
- ☐ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$690.00
- ☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$670.00
- ☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfy provisions of PCT Article 33(1)-(4) \$96.00

CALCULATIONS PTO USE ONLY**ENTER APPROPRIATE BASIC FEE AMOUNT =****\$840.00**

Surcharge of **\$130.00** for furnishing the oath or declaration later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).

\$0.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	
Total claims	31 - 20 =	11	x \$18.00	\$198.00
Independent claims	4 - 3 =	1	x \$78.00	\$78.00
Multiple Dependent Claims (check if applicable).			<input type="checkbox"/>	\$0.00

TOTAL OF ABOVE CALCULATIONS = \$1,116.00

Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28) (check if applicable). ☐

\$0.00**SUBTOTAL = \$1,116.00**

Processing fee of **\$130.00** for furnishing the English translation later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).

\$0.00**TOTAL NATIONAL FEE = \$1,116.00**

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable). ☐

\$0.00**TOTAL FEES ENCLOSED = \$1,116.00**

Amount to be refunded	\$
charged	\$

- ☒ A check in the amount of **\$1,116.00** to cover the above fees is enclosed.
- ☐ Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.
- ☒ The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account No. **19-2253** A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

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**24223**

PATENT TIMELINE OFFICE

SIGNATURE

Michael I. Stewart

NAME

24,973

REGISTRATION NUMBER

August 22, 2000

DATE

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Our Ref: 6900-255 MIS:jb

In re National Phase of International Application

No.: PCT/CA99/00155

International

Filing Date: 22 February 1999

Applicant: Zhi-Zhong Liang

Title: LIGNIN BASED COLLOIDAL COMPOSITIONS

PRELIMINARY AMENDMENT

The Commissioner of Patents
and Trademarks,
Washington, D.C. 20231,
U. S. A.

Dear Sir:

Please amend this application in the following manner:

In the Disclosure:

Before the first line of the specification, add the following:

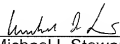
" REFERENCE TO RELATED APPLICATIONS

This application is a national phase application under 35 U.S.C. 371 of
PCT/CA99/00155."

REMARKS

The specification has been amended on page 1 to reflect that this
application is a U.S. National Phase filing under 35 U.S.C. 371 of PCT/CA99/00155.

Respectfully submitted,



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Date: August 22, 2000

000211-1322960

PTO/PCT Rec'd 23 AUG 2000

TITLE OF INVENTIONLIGNIN BASED COLLOIDAL COMPOSITIONSFIELD OF INVENTION

5 The present invention relates to the processing of
black liquor and the preparation of bitumen compositions
containing black liquor.

BACKGROUND TO THE INVENTION

10 Black liquor is an aqueous alkaline pulping residue
produced in Kraft pulping of wood chips in a pulp mill
to provide wood fibers for paper-making. Black liquor
generally contains about 55 to 65% solids comprising
lignin, sodium salts of poly-saccharinic acids, organic
and inorganic sulfides and carbonates and many other
compounds. The lignin in the black liquor is the main
15 organic component and is present as a high mass colloid
stabilized by ionized phenolic groups. Lignin is a
phenylpropane polymer of amorphous structure comprising
about 17 to about 30% of wood. Lignin is so closely
associated with the cellulosic fibres and hemi-cellulose
20 that makes up the balance of woody material that lignin
can be separated from wood only by chemical reaction at
high temperature, such as that used in the Kraft pulping
process, wherein wood chips are digested with an aqueous
alkaline solution of sodium hydroxide and hydrogen
25 sulfide, known as white liquor. Normally, the black
liquor, after separation of the wood fibers, is
processed to remove organic materials and to recover and
regenerate pulping chemicals. Lignin can be recovered
from wood-processing wastes only in limited quantities.

30 The lignin is a natural binder derivative of wood.
Alkali lignin from both softwoods and hardwoods are
useful in many products. They are used as stabilizers
for asphalt emulsions, modifiers and extenders for latex
emulsions, compounding agents in vinyl plastics, soil
35 conditioners, binders in printing inks, wood stains,
protective colloids in soap emulsions, dispersing agents

for clay, fire foam stabilizers, drilling mud additives, insecticide dispersants, absorbers in storage batteries and foundry sand binders. Alkali lignin can also be used as adhesives for plywood, molding powders and formed insulating resins and as a reinforcing agent for rubber.

Asphalt binders (or bitumen) are widely used in different applications, such as asphalt-aggregate blends for road paving, asphalt fiber reinforced membranes for roofing and asphalt-water emulsions in surface treating both for paving and roofing. Problems for straight-run asphalt in those applications are the tendency to become brittle at low temperatures and to become soft at high temperatures, the tendency to flow under constant load at service temperatures and the tendency to phase separation during storage when mixed with other components.

Various approaches have been used for the purpose of improving asphalt compositions to address these problems. For instance, polymers have been utilized to improve the high and low temperature characteristics of asphalt compositions, as well as to improve their toughness and durability. Air blowing of straight-run asphalt has been used to improve stiffness and dimensional stability for hot applied roofing systems. The asphalt may be treated with acid to improve its compatibility with the polymeric additives. Lignin or lignin based additives have been widely used to emulsify asphalt to achieve a stable dispersion of asphalt in aqueous medium.

Black liquor is an aqueous colloidal system in which ample lignin is present. However, means have not been previously disclosed whereby a stable dispersion of lignin in asphalt can be achieved by combining black liquor with hot liquid asphalt. When the black liquor is added to the bitumen and dispersed with agitation into molten asphalt cement, de-watering of the black liquor

It is one objective of the present invention to develop a process or processes whereby liquor-solid pulping residue (black liquor) can be converted from one in which water is the dispersing and plasticizing agent to one in which water is replaced by oil, achieving a dispersion in which a lignin based colloidal particulate phase is stably maintained. It is another objective to provide a lignin-oil dispersion which can be smoothly blended with liquid asphalt to produce a stable, homogeneous dispersion.

The present invention relates to compositions and methods for combining black liquor with selected petroleum-derived products. More specifically, this invention provides two separate processes for preparing lignin-based colloidal dispersions in lubricating or other petroleum-based oils. The lignin-oil colloidal dispersions prepared by either of such processes may be blended with bitumen to produce novel bituminous composition. Two embodiments of such procedures are described herein. The black liquor used in the invention may be that resulting from cooking wood in an alkaline solution in the soda or sulfate (Kraft) paper-making process.

In a first embodiment of the invention, the

procedure involves essentially two steps. In the first step, the black liquor is converted from its aqueous colloidal system to an anhydrous colloidal dispersion in a lubricating oil medium in which a surfactant, which may be an anionic surfactant, such as DDBSA (dodecylbenzene sulfuric acid), is used to promote the provision of the anhydrous colloidal dispersion. The process may be effected under low shear conditions.

The anhydrous colloidal dispersion of lignin in oil may be mixed, in a second step, with asphalt bitumen, producing a unique combination, which is a novel bituminous composition.

In another embodiment of the invention, the procedure again involves essentially two steps. First, bitumen is treated with an inorganic acid, such as sulfuric acid, at elevated temperature to provide an acid-treated bitumen. The black alkaline liquor is then added slowly to the acid-treated bitumen under agitation for de-watering the black liquor at a controlled rate to form a stable, substantially anhydrous, colloidal dispersion of lignin in the bitumen, also to provide a novel bituminous composition.

Accordingly, in another aspect of the present invention, there is provided a bituminous composition comprising bitumen, and dewatered lignin-containing spent alkaline pulping liquor dispersed therein. The dewatered spent alkaline pulping liquor may be present in an amount up to about 50 wt.% of the bitumen, preferably about 5 to about 25 wt.% of the bitumen.

In such composition, the dewatered spent alkaline pulping liquor may comprise an anhydrous colloidal dispersion of lignin in a lubricating oil. The dewatered spent alkaline pulping liquor may comprise a stable, substantially anhydrous, colloidal dispersion of lignin in the bitumen.

DETAILED DESCRIPTION OF THE INVENTION

Referring first to the first embodiment of the invention, a lubricating oil is mixed with black liquor at temperatures below the water boiling point. A
5 surfactant, which may be DBSA (dodecylbenzene sulfuric acid), is selected to be added to the mixture to facilitate the colloidal dispersion of black liquor in the lubricating oil while de-watering is gradually carried out at an elevated temperature. The dewatering
10 may be effected by starting at an elevated temperature, which may be about 100° to about 180°C, preferably about 110° to about 160°C. The resulting cream-like paste product is miscible with hot liquid asphalt at any ratio to form a novel bituminous composition.

15 The lubricating oil used in the first embodiment of the invention may be a re-refined waste motor oil or other convenient petroleum-based oil. Contaminants of waste motor oil, such as dirt, lead, arsenic and other harmful metals and chemicals, are removed from the waste
20 motor oil to produce a clean base lubricating oil by the re-refining process.

The lubricating oil may be a selected fraction of refined mineral oil used for lubrication of moving surfaces. Such fraction may range in consistency from
25 thin liquid to grease-like substances. Usually, lubricating oils contain small amounts of additives to impart special properties, such as viscosity index and detergency.

The surfactant or dispersing agent used in the
30 process may typically be dodecylbenzene sulfuric acid (DBSA), although other anionic surfactants may be employed, if desired, such as fatty acid, linear alkyl sulfonates having 10 or more carbon atoms in the chain. DBSA is benzene with dodecene, and the resulting
35 dodecylbenzene is sulfonated. DBSA may be neutralized with caustic soda from the alkaline black liquor or

other convenient source to promote the detergency of the lubricating oil and to reduce surface tension in the mixture.

5 However, a surfactant need not always be necessary, if the lubricating oil employed has a detergency high enough to ensure that a colloidal dispersion of the solid residue from the black liquor is achieved in the lubricating oil medium.

10 The lignin-oil dispersion produced by either of these procedures may be smoothly blended with bitumen to provide the desired lignin-asphalt composition. The spent pulping liquor, which may be black liquor from a kraft pulp mill, may be present in an amount of about 25 to about 60 wt.% of the composition, preferably about 25 to 40 wt.% of the composition.

15 Referring now to the second embodiment of the invention, the bitumen first is treated with a strong mineral acid, such as sulfuric acid, at elevated temperature and then the alkaline black liquor is added to the treated bitumen at a controlled rate under agitation to effect de-watering of the black liquor to form a stable, substantially anhydrous, colloidal dispersion in the bitumen. The elevated temperature may be about 100° to about 180°C, preferably about 110° to about 160°C. The spent pulping liquor may be present in an amount of about 5 to about 50 wt.% of the composition, preferably about 10 to about 30 wt.%

20 An inorganic acid is contacted with, or added to, the asphalt to form an acid treated asphalt. In general, 30 the acid addition shifts the asphalt structure from a sol to a gel, lowers the temperature susceptibility of the asphalt and improves the stability of the additive dispersion in the treated asphalt. Since black liquor is an aqueous alkaline system, it is critical that the acid not be added to the asphalt after addition of or 35 with the black liquor.

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Preferably, the acid is added slowly to the asphalt to avoid foaming, which may occur if all the acid were added at one time. The inorganic acid content of the asphalt resulting from the acid treatment is not critical, but normally is in the range between about 0.2 and about 3.5 wt%, preferably between about 0.5 and about 2.5 wt% of the asphalt. Although a wide variety of inorganic acids can be used for treatment of the asphalt, the inorganic acid is preferably selected from the group consisting of sulfuric acid, phosphoric acid, poly-phosphoric acid, phosphorous pentoxide, hydrochloric acid, and mixtures thereof. The sulfuric acid, phosphoric acid or poly-phosphoric acid are preferred inorganic acids, with the sulfuric acid being particularly preferred.

Following acid addition, the alkaline black liquor is then added slowly to the acid-treated bitumen at an elevated temperature, typically around 100°C, under agitation for a certain period of time to permit de-watering of the black liquor to occur while the solids are incorporated into the bitumen. The temperature of the final composition next is increased above the boiling point of water, typically up to around 160°C until residual water is completely evaporated off to

form a stable, substantially anhydrous, colloidal lignin dispersion in the bitumen.

EXAMPLES

Example 1:

- 5 This Example illustrates the first embodiment of the invention in which black liquor first dispersed in a lubricating oil and then in bitumen.

General Procedure:

- 10 Black liquor and DDBSA were mixed at room temperature in specific proportions under stirring for 20 to 30 min to form a mixture. Re-refined waste lube-oil was added into the mixture at room temperature to a desired concentration and stirred another about 20 min. The resulting mixture was heated slowly to about 160° to 15 180°C for about 2 to 3 hours under stirring until water-free. During this step, the viscosity of the system exhibited complex phenomena because a raising of the temperature resulted in reduced viscosity while water loss resulted in increased viscosity. Accordingly, the 20 temperature and time for de-watering must be controlled.

Specific Experiments:

- 16 Kg of DDBSA was added to 266 g of black liquor (ca. 60% solid content) and stirred at room temperature for 20 to 30 min. 184 g of reclaimed lube oil were added 25 and stirred again at room temperature for 20 to 30 min. The temperature of the mixture was slowly raised from room temperature to 170°C for 2 to 3 hours to de-water the composition. After de-watering and cooling, about 332 g sample was obtained. The sample was a black 30 colored paste and water-free and was easily dispersed into asphalt at any ratio. This sample was designated as B-14, seen in Table 1.

- Different black liquor/lube oil formulations have been prepared following the above procedure and the data 35 are presented in the following Table 1:

TABLE 1
NON-AQUEOUS DISPERSING SYSTEM OF BLACK LIQUOR/LUBE OIL

Sample Code	Black liquor g	DBSA		Lube Oil		Product Wt. g
		g	wt%	g	wt%	
B-13	246	24	4.6	254	48.5	410
B-14	266	16	3.4	184	39.5	332
B-15	218	6.0	1.94	180	44.6	302
B-19	336	10	2.15	120	25.8	308
B-20	460	14	2.35	122	20.5	372
RB-30	590 (1 part)	17.7 (0.03 part)		413 (0.7 part)		

Note: Mass of analyzed sample of reclaimed lube oil (recovered from engine oil) was determined following the test method of ASTM 4124-92.

Base asphalt (or bitumen) used in the present invention were selected from three different degrades: AC-5 having an absolute viscosity of around 500 poises, I85/100 having a 25°C penetration range from 85 dmm to 100 dmm and I120/150 having a 25°C penetration range from 120 dmm to 150 dmm. The non-aqueous dispersing system of black liquor/lube-oil produced as described in the Table 1 was dispersed into hot liquid asphalt at about 160°C respectively to form a very fine dispersion mixture. The stability of these lignin based bituminous mixtures during hot storage at elevated temperature was evaluated using a conditioned tube test. The conditioning procedure consists of placing approximately 70g of the mixture in ¾" aluminum tubes and storing such tubes in a vertical position at 165°C in an oven for 48 hrs. Following hot storage, a viscosity ratio was determined by comparing the viscosity of the mixture tested at 135°C from the bottom section of the sample and from the top section of the sample in the tube. A ratio in the range of 0.80 to 1.20 is generally considered acceptable with respect to separation of the dispersed phase. The testing results on eight compositions are shown in the following Table 2. The results indicate that the lignin based solid dispersion in asphalt binder are quite stable.

TABLE 2
STABILITY OF THE BLACK LIQUOR/LUBE OIL BLEND (BLE) IN HOT ASPHALT

Code	Asphalt		BLB		Viscosity, cp at 135°C	Stability Index*
	Name	(g)	Code	(g)	Top/Bottom	
AB5-16	ACS	368	B-13	38	455/463	0.98
B14-85-10	I85/100	90	B-14	10	387.5/432.5	0.90
B14-85-20	I85/100	80	B-14	20	450/465	0.97
B14-85-30	I85/100	70	B-14	30	435/500	0.87
B15-A	I120/150	70	B-15	30	825/845	0.97
B15-B	I120/150	85	B-15	15	257/265	0.97
B20-20	I120/150	80	B-20	20	740/802.5	0.92
RB30-30	I120/150	80	RB-30	20	405/475	0.85

Note: (1) In the better formulations of the non-aqueous dispersing system, an appropriate composition was a lube oil content of more than 20% and DBSA content of greater than 2%.

(2) DBSA is dodecylbenzene sulfonic acid.

Stability Index = $\frac{\text{Viscosity at top}}{\text{Viscosity at bottom}}$

Example 2:

This Example illustrates the second embodiment of the invention in which bitumen first is acidified.

General Procedure:

5 Sulfuric acid (95 to 98% H_2SO_4) was added into hot asphalt AC-5 at 100° to 110°C under stirring (800 to 1000 RPM) for 20 to 30 mins. to form an acid-modified asphalt, followed by addition of aqueous black liquor (c.a. 60% solid content) into the acid-modified asphalt, according to the following procedure.

- (1) While stirring at 800 to 1000 RPM, the acid was added into asphalt at 100° to 110°C for 20 to 30 min. and acid-modified asphalt was obtained.
- 15 (2) After 20 to 30 min., black liquor was added to the acid-modified asphalt at 105° to 115°C for 30 to 40 min., then maintained for 1 to 2 hours for dewatering.
- (3) The temperature was slowly increased to 170°C while stirring for 1 to 2 hours to effect further dewatering.
- 20 (4) The composition was allowed to cool and was sampled for stability.

Specific Experiments:

25 To 392 g of asphalt AC-5 were added 3.14 g of H_2SO_4 (0.8% AC-5) at 100° to 110°C under 800 to 1000 RPM stirring for 20 min. The acidified asphalt was maintained at 100° to 110°C and black liquor 84 g (22% AC-5) was added over about 30 min., and then the composition was maintained for 1 to 2 hrs at 105° to 30 115°C to dewater the black liquor. The temperature was raised to 170°C while stirring to effect further dewatering. The product was then cooled to provide 436 g of product (loss mass 43 g). The stability index determined on this sample (AC-B-18) was 0.94, shown in 35 Table 3.

Results of dispersion and stability determinations are also shown below in the following Table 3. The results shown in Table 3 indicate that products of formulation containing 0.8 wt% acid and 0 to 30 wt%, preferably 15 to 25 wt% black liquor, based on the amount of asphalt, had good stability.

TABLE 3
EXPERIMENTAL FORMULATIONS AND STABILITY

Code	Asphalt (AC-5)	Acid		Black Liquor		Viscosity cp after 48 hrs storage at 160°C		
		(g)	(%)	(g)	(%)	Top	Bottom	Stability Index
AC-B-24	374	0	0	112	30			
AC-B-11	472	2.4	0.5	94	20	413	550	0.75
AC-B-10	362	4.0	1.1	72	20	298	310	0.96
AC-B-18	392	3.14	0.8	84	22	338	358	0.94
AC-B-25	294	2.35	0.8	59	20	312.5	317.5	0.98
AC-B-26	288	2.3	0.8	72	25	360	342.5	1.05
AC-B-19	360	2.94	0.8	110	30	270	598	0.45
AC-B-23	380	3.8	1.0	114	30	290	907.5	0.32
AC-B-21	360	4.32	1.2	108	30	377.5	360	1.05
AC-B-22	373	3.73	1.0	149	40	300	402.5	0.75
AC-B-20	380	3.0	0.8	190	50	267.5	1060	0.25

Note: (1) No measurement was possible.

SUMMARY OF THE DISCLOSURE

In summary of this disclosure, the present invention provides novel procedures for the stable dispersion of lignin components of black liquor in bitumen. Modifications are possible within the scope of the invention.

5

000211-18222960

CLAIMS

What I claim is:

1. A bituminous composition, comprising
bitumen, and
dewatered lignin-containing spent alkaline pulping liquor stably dispersed therein.
2. The bituminous composition of claim 1 wherein said dewatered spent alkaline pulping liquor is present in an amount up to about 50 wt% of the bitumen.
3. The bituminous composition of claim 2 wherein said dewatered spent alkaline pulping liquor is present in an amount of from about 5 to about 25 wt% of the bitumen.
4. The bituminous composition of claim 1 wherein said dewatered spent alkaline pulping liquor comprises an anhydrous colloidal dispersion of lignin in a lubricating oil.
5. The bituminous composition of claim 1 wherein said dewatered spent alkaline pulping liquor comprises a stable, substantially anhydrous, colloidal dispersion of lignin in the bitumen.
6. The bituminous composition of claim 1 wherein the spent alkaline pulping liquor is black liquor from a Kraft pulp mill.
7. A method of producing a composition, which comprises:
converting an aqueous colloidal dispersion of liquor in spent pulping liquor to an anhydrous colloidal dispersion of lignin in a lubricating oil.
8. The method of claim 7 wherein said converting step is effected by:
mixing a lubricating oil with the spent pulping liquor, and
dewatering the mixture so formed at an elevated temperature to form a cream-like paste compatible with hot bitumen.

9. The method of claim 8 wherein said mixing step is effected in the presence of a surfactant to facilitate colloidal dispersion of the spent pulping liquor in the lubricating oil.
10. The method of claim 9 wherein said surfactant is an anionic surfactant.
11. The method of claim 10 wherein the anionic surfactant is a fatty acid linear alkyl sulfonate.
12. The method of claim 7 including blending the anhydrous colloidal dispersion of lignin in a lubricating oil with bitumen.
13. The method of claim 7 wherein said spent pulping liquor is black liquor from a Kraft pulping mill.
14. The method of claim 7 wherein the spent pulping liquor is present in an amount of about 10 to about 60 wt% of the composition.
15. The method of claim 14 wherein the black liquor is present in an amount of about 25 to 40 wt%.
16. A method of producing a bituminous composition, which comprises:
- converting an aqueous colloidal dispersion of lignin in spent alkaline pulping liquor to an anhydrous colloidal dispersion of lignin in a lubricating oil, and
- mixing said anhydrous colloidal dispersion of lignin with bitumen.
17. The method of claim 16 wherein said spent alkaline pulping liquor is black liquor from a Kraft pulp mill.
18. The method of claim 16 which is effected by stressing at elevated temperature.
19. The method of claim 18 wherein said elevated temperature is about 100° to about 180°C.
20. The method of claim 19 wherein said temperature is about 110° to about 160°C
21. A method of producing a bituminous composition which comprises:
- treating bitumen with an inorganic acid at an

elevated temperature to provide acid-treated bitumen,

slowly adding to the acid-treated bitumen a lignin-containing spent alkaline pulping liquor, while agitating the bitumen to effect controlled dewatering of the spent pulping liquor to form a stable, substantially anhydrous colloidal dispersion of lignin in the bitumen.

22. The method of claim 21 wherein said inorganic acid treatment of bitumen is effected to convert the bitumen structure from a sol to a gel, lower the temperature susceptibility of the asphalt and improve the stability of the additive dispersion in the treated bitumen.

23. The method of claim 22 wherein said inorganic acid is added slowly to the bitumen at an elevated temperature to avoid foaming to provide an inorganic acid content of said bitumen of about 0.2 to about 3.5 wt%.

24. The method of claim 23 wherein said inorganic acid content is from about 0.5 to about 2.5 wt% of the bitumen.

25. The method of claim 21 wherein said inorganic acid is sulfuric acid.

26. The method of claim 21 wherein said addition step is effected by:

initially slowly adding the spent alkaline pulping liquor to the acid-treated bitumen at substantially water boiling temperature while agitating the bitumen to effect an initial dewatering of the alkaline spent pulping liquor, and

subsequently increasing the temperature of the bitumen above the water boiling point to effect a further dewatering of the bitumen to form the stable, substantially anhydrous, colloidal dispersion of lignin in the bitumen.

27. The method of claim 21 wherein said spent alkaline pulping liquor is black liquor from a Kraft pulp mill operation.

28. The method of claim 21 wherein said elevated temperature is about 100° to about 180°C.

29. The method of claim 28 wherein said temperature is about 110° to about 160°C.

30. The method of claim 21 wherein the spent pulping liquor is present in an amount of about 5 to about 50 wt% of the composition.

31. The method of claim 21 wherein the spent pulping liquor is present in an amount of about 10 to about 30 wt%.

Docket No.
6900-255 MIS

Declaration and Power of Attorney For Patent Application

English Language Declaration



As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

LIGNIN BASED COLLOIDAL COMPOSITIONS

the specification of which

(check one)

☐ is attached hereto.

☒ was filed on 22 February 1999 as United States Application No. or PCT International Application Number PCT/CA99/00155

and was amended on _____

(if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose to the United States Patent and Trademark Office all information known to me to be material to patentability as defined in Title 37, Code of Federal Regulations, Section 1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, Section 119(a)-(d) or Section 365(b) of any foreign application(s) for patent or inventor's certificate, or Section 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate or PCT International application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)

Priority Not Claimed

_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/>
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/>
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/>

I hereby claim the benefit under 35 U.S.C. Section 119(e) of any United States provisional application(s) listed below:

60/075,546

23 February 1998

(Application Serial No.)

(Filing Date)

(Application Serial No.)

(Filing Date)

(Application Serial No.)

(Filing Date)

I hereby claim the benefit under 35 U. S. C. Section 120 of any United States application(s), or Section 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. Section 112, I acknowledge the duty to disclose to the United States Patent and Trademark Office all information known to me to be material to patentability as defined in Title 37, C. F. R., Section 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application:

PCT/CA99/00155

22 February 1999

(Application Serial No.)

(Filing Date)

(Status)
(patented, pending, abandoned)

(Application Serial No.)

(Filing Date)

(Status)
(patented, pending, abandoned)

(Application Serial No.)

(Filing Date)

(Status)
(patented, pending, abandoned)

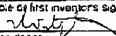
I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

- POWER OF ATTORNEY. As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith. (list name and registration number)

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